Prisca 5.1.0.17

Date of report: 16/08/21

SONIYA

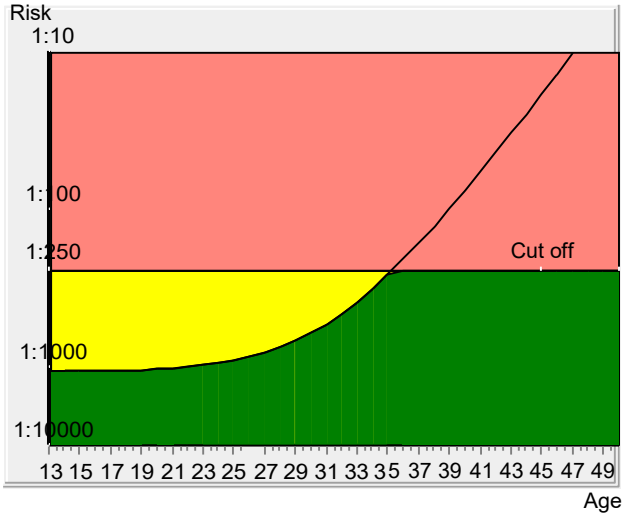
Patient data			
Name	Mrs. VANI	Patient ID	0012108160308
Birthday	12/05/95	Sample ID	22501624
Age at sample date	26.3	Sample Date	16/08/21
Gestational age	11 + 3		

Correction factors					
Fetuses	1	IVF	no	Previous trisomy 21 pregnancies	unknown
Weight	52.6	diabetes	no		
Smoker	no	Origin	Asian		

Biochemical data		
Parameter	Value	Corr. MoM
PAPP-A	4.7 mIU/n L	1.66
fb-hCG	56.89 ng/mL	1.03

Risks at sampling date	
Age risk	1:865
Biochemical T21 risk	<1:10000
Combined trisomy 21 risk	<1:10000
Trisomy 13/18 + NT	<1:10000

Ultrasound data	
Gestational age	11 + 3
Method	CRL Robinson
Scan date	16/08/21
Crown rump length in mm	49
Nuchal translucency MoM	0.90
Nasal bone	unknown
Sonographer	DR NA
Qualifications in measuring NT	MD



Trisomy 21

The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.

After the result of the Trisomy 21 test (with NT) it is expected that among more than 10000 women with the same data, there is one woman with a trisomy 21 pregnancy.

The calculated risk by PRISCA depends on the accuracy of the information provided by the referring physician.

Please note that risk calculations are statistical approaches and have no diagnostic value!

The patient combined risk presumes the NT measurement was done according to accepted guidelines (Prenat Diagn 18: 511-523 (1998)).

The laboratory can not be hold responsible for their impact on the risk assessment ! Calculated risks have no diagnostic value!

Trisomy 13/18 + NT

The calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:10000, which represents a low risk.

Sign of Physician

below cut off | Below Cut Off, but above Age Risk | above cut off